

GUIDED-MODE LASER APPARATUS WITH IMPROVED CLADDING STRUCTURE AND
A METHOD OF FABRICATING THEREOF

I. BACKGROUND OF THE INVENTION

1. Field of the Invention.

This invention relates to the field of crystalline materials to be used in fabrication of the core and the cladding of diffusion bonded optics. More particularly, it pertains to the use of undoped yttrium aluminum garnet (hereinafter, YAG) for the fabrication of the cladding, and of doped lutetium aluminum garnet (hereinafter, LuAG) for the fabrication of the core, the core being a rod completely surrounded by the cladding on all sides including the ends of the rod.

2. Description of the Related Art.

Many modern waveguided structures comprise a core diffusion-bonded to cladding in which this core is completely ensconced (except for the ends). Prior art for such structures incorporated sapphire as the cladding material. For example, U.S. Patent No. 5,852,622 to Meissner et. al., teaches a system for improving the

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Laser guiding, via total internal reflection (TIR), takes place when the refractive index of the core is larger than that of the cladding. TIR occurs when the angle of incidence is larger than the critical angle θ_{critical} , where θ_{critical} is determined from the equation $\sin \theta_{\text{critical}} = n_{\text{cladding}}/n_{\text{core}}$, where n_{cladding} and n_{core} are refractive indices of the cladding and the core, respectively.

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It follows from the foregoing that the angular spread of what is guided depends on the difference in the refractive indices, and the advantages of sapphire include its much lower refractive index relative to YAG (1.76 and 1.82, respectively).

In addition, sapphire has superior thermal conductivity compared to that of YAG (about 35 Watts per meter per Kelvin for sapphire compared to 10 Watts per meter per Kelvin for YAG), which is important to facilitate the removal of heat from the YAG core.

However, this prior art design employing sapphire had serious disadvantages, particularly in that it caused difficulties with the six sided bonding using sapphire. The dissimilar crystalline and mechanical properties make the practical fabrication of diffusion-bonded fully encased sapphire and YAG composite structures very difficult. Those skilled in the art are aware

that the manufacturing of diffusion-bonded ytterbium-doped YAG-sapphire composites, promoted by Onyx Optics, Inc. of Dublin, California, albeit successful, has proved to be very time-consuming as well as very expensive undertaking.

Specifically, because the polishing rate for the two materials is different, it is very difficult to obtain an adequately flat optical finish over a sapphire-YAG-sapphire composite surface.

Another drawback of the sapphire-based technology is that according to this technology it is essential to obtain surface free of scratches, gaps, voids, inclusions, digs and similar mechanical imperfections in preparing the surface for any subsequent bonding steps. Although this disadvantage may characterize any known diffusion bonding technique, in the case of YAG it is mitigated because it is easier to polish YAG than to polish sapphire.

Also, the crystalline axis of the sapphire (a uniaxial crystal) must be oriented in a particular direction relative to the YAG interface for reliable diffusion bonding. A simple geometrical analysis shows that this preferred orientation is not possible for bonding on all six sides. Hence in practice, a YAG core fully

A specific embodiment of the present invention uses undoped YAG as the material for making the cladding and uses ytterbium or neodymium-doped LuAG as the material for the fabrication of the core, the core being completely surrounded by the cladding on all sides including the ends of the rod.

LuAG has a slightly higher refractive index than YAG, so optical guiding occurs. The Yb-doped LuAG has the same general absorption and emission characteristics as the prior-art Yb-doped YAG core. Because both YAG and LuAG are aluminum garnets, they are quite similar in physical, optical, and thermomechanical properties. Hence, it is significantly easier to achieve diffusion-bonding of this composite than the bonding of YAG to sapphire described in prior art. Use of these materials for the core and the cladding makes diffusion-bonding fabrication processes significantly easier while maintaining adequate, waveguiding and spectral characteristics.

This invention provides the following improvements over the prior art.

1. By using a cladding material, physical and thermomechanical properties of which, such as linear-thermal expansion, are nearly ideally matched to the core material, this invention allows much

threshold is 30% lower.

Therefore it is easier to extract energy from ytterbium-doped LUAG core for a given set of parameters compared to ytterbium-doped YAG core used in prior art. Other key spectral parameters are also maintained on good levels when the materials of this invention are used to fabricate the device. For instance, the absorption cross-section of the ytterbium-doped LUAG core is comparable to that of the ytterbium-doped YAG core used in prior art. The fluorescence lifetimes of the two materials are also very close at 925 μ s and 951 μ s, respectively.

5. The thermal expansion coefficient of LuAG is slightly larger than that of YAG. Therefore, under thermal load, LuAG is under compressive stress, which means the design is more thermally robust than if the core were under tensile stress.

Other particular embodiments of this invention may benefit by any or all of these improvements depending on the specific selection of lasing medium and cladding material. Criteria for such selection and the materials selected are discussed subsequently.

One aspect of this invention provides a solid state waveguided

structure comprising a core fabricated of a lasing medium, the core having an outer surface, and a cladding fabricated of a laser-inactive material, said cladding diffusion-bonded to the outer surface of the core, wherein the lasing medium of the core comprises a crystalline or a glass material, preferably, a lutetium-aluminum-garnet material doped with ions, preferably, with ions of ytterbium, lutetium-aluminum-garnet material doped with ions of neodymium, and lutetium-aluminum-garnet material doped with ions of thulium, and the laser-inactive material of the cladding comprises an yttrium-aluminum-garnet material.

Another aspect of this invention provides a method of fabricating a solid state waveguided structure with improved characteristics, comprising steps of providing a core fabricated of a lasing medium, the core having an outer surface, and ensconcing the core in a cladding fabricated of a laser-inactive material, the cladding diffusion-bonded to the outer surface of the core, wherein the lasing medium of the core comprises a crystalline or a glass material, preferably, a lutetium-aluminum-garnet material doped with ions, preferably, ions of ytterbium, lutetium-aluminum-garnet material doped with ions of neodymium, and lutetium-aluminum-garnet material doped with ions of thulium, and the laser-inactive material of the cladding comprises an yttrium-

aluminum-garnet material.

III. BRIEF DESCRIPTION OF THE DRAWINGS

The features and advantages of the present invention will become better understood with regard to the following description, appended claims, and accompanying drawings where

FIG. 1a is a schematic diagram showing a preferred architectural structure of the composite device of this invention.

FIG. 1b is a schematic diagram showing an end view of the structure depicted on FIG. 1a, when looked at in direction A.

FIG. 2 is a chart showing a results of thermal analysis of various active cores.

IV. DETAILED DESCRIPTION OF THE INVENTION

In accordance with the present invention, a guided wave amplifier which maintains the gain achievable at the ytterbium-doped YAG signal wavelength, as known in prior art, but in a different host crystalline medium, is provided. Using different crystalline

medium, described above, facilitates the manufacturing process and lowers the cost.

FIG. 1a shows a preferred architectural structure of the composite device of this invention, and FIG. 1b depicts an end view of the structure shown on FIG. 1a, when looked at in direction A.

The device 100 comprises a preferably cylindrical central section 1 and two substantially identical end sections 2a and 2b, which are also preferably cylindrical.

Each of these end sections 2a and 2b has an inner side 3a and 3b, respectively, and an end face 4a and 4b, respectively. End faces 4a and 4b are normally fabricated at an angle away from normal to the core rod axis; this angle is calculated according to methods known to those skilled in the art and is large enough that Fresnel reflections from the air/crystal interface cannot couple into the core 5. It is usual but not essential that the ends of the core 5 be normal to the rod axis.

Between the central section 1 and each of the end sections 2a and 2b, there lie two substantially identical "bottleneck" sections

7a and 7b, respectively. Each "bottleneck" section 7a and 7b has a diameter which progressively increases, from its lowest value at the point 7c (where the "bottleneck" section 7a or 7b touches the central section 1) to its highest value at the point 7d (where the "bottleneck" section 7a or 7b merges into the end section 2a or 2b, respectively).

Each of the inner faces 3a and 3b is preferably circular in cross-section and is diffusion-bonded to the central section 1 so that the central section 1 is integral with said the end sections 2a and 2b. Each of the end faces 4a and 4b is preferably circular or elliptical.

The central section 1 is comprised of a core 5, said core 5 being completely ensconced in cladding 6. The core 5 is a rod, polygonal in cross-section, preferably a square, as shown on FIG. 1b. Those skilled in the art will realize, that depending on the refractive index difference between the cladding and the core, various polygonal forms can be used.

This rod, forming the core 5, therefore, in the preferred embodiment, preferably has four sides and two ends, a total of six surfaces, and extends lengthwise throughout the central section 1.

Cladding 6 comprises both the cladding's portion 6a of which end sections 2a and 2b are made and the cladding's portion 6b, enveloping the core 5 in the central section 1.

The core 5 is fabricated of crystalline or a glass material, preferably, LuAG, doped with the trivalent ion of ytterbium, Yb^{3+} , and the cladding 6 is preferably fabricated of clear undoped YAG.

Therefore, the preferred embodiment of this invention comprises a device having the ytterbium-doped LUAG core and YAG cladding. Those skilled in the art will realize that other combinations of the core and of the cladding materials can be chosen. The criteria for the selection of the materials are as follows:

1. The properties of the materials of the core and of the cladding should match.
2. Such end caps should be used that the beam size at the air-crystal interface is larger than the size in the core.
3. The doped core should be made of a material having slightly higher refractive index than the material of the cladding.

4. Spectral characteristics of the materials should be appropriately chosen.

5. Thermal expansion of the material of the core should be slightly higher than that of the material of the cladding.

Using the foregoing criteria, selection of materials for the core and for the cladding may lead to a choice of materials comprising a neodymium-doped LUAG core/YAG cladding combination or a thulium-doped LUAG core/YAG cladding combination, in addition to the ytterbium-doped LUAG core/YAG cladding combination. Garnets other than YAG can be used, but YAG is a preferred garnet in terms of combination of properties as well as availability.

The garnet used in the preferred embodiment of the invention is similar to YAG used in prior art, but its use does not bring about any drawbacks and disadvantages inherent in YAG which were discussed hereinabove. However, same pump diodes which are used in ytterbium-doped YAG cores of prior art can be used with ytterbium-doped LUAG core of this invention and the gain achieved at ytterbium-doped YAG wavelength is similar as well.

The preferred embodiment of this invention is depicted on FIGs. 1a

The composition of the preferred embodiment, ytterbium Yb^{3+} -doped LuAG material for the core 5 and clear undoped YAG for the cladding 6, satisfies the requirements of the thermal and mechanical properties of the materials of the cladding 6 and of the core 5 discussed above. In addition, the refractive indices of LuAG and YAG are close, LuAG having somewhat higher refractive index, so that guiding can occur with LuAG as the core medium. The resulting numerical aperture (NA) is calculated according to a formula $(n_{\text{core}}^2 - n_{\text{cladding}}^2)^{1/2}$ or $(1.83^2 - 1.82^2)^{1/2}$ and is about 0.19, which is an excellent number for NA.

The relevant thermomechanical properties as well as refractive indices for YAG and LuAG are summarized in Table 1. The comparison data for sapphire are also given.

TABLE 1.

COMPARISON OF THERMOMECHANICAL PROPERTIES AND REFRACTIVE INDICES OF YAG AND LUAG

Material	Thermal Conductivity, W/m/K	Young Modulus E, GPa	Thermal Linear Expansion Coefficient (1/K)	Refractive Index, n_D , at 1 μm
Sapphire	~35	405	5.4x10 ⁻⁶ //a axis 6.8x10 ⁻⁶ //c axis	1.76
YAG	10	282	7.0x10 ⁻⁶	1.82
LuAG	9	300	8.8x10 ⁻⁶	1.83

In order to maintain a reasonable absolute temperature, T, in the LuAG core, dimensions of the portions of this device are very important. The set of the acceptable and the preferred dimensions is summarized in Table 2.

TABLE 2.

TYPICAL DIMENSIONS OF PORTIONS OF THE STRUCTURE OF THIS INVENTION
(PREFERRED EMBODIMENT)

Portion of the Device	Dimension shown as on FIGs. 1a or 1b	Dimension, within a range of	Dimension, preferred
Central section 1	"a" on FIG. 1a	Between about 23 and about 25 mm	24 mm
Between point 7d of the end of the "bottleneck" section 7a or 7b and point 7e*	"b" on FIG. 1a	Between about 2.4 and about 2.6 mm	2.5 mm
Between point 7c of the beginning of the "bottleneck" section 7a or 7b and point 7e*	"c" on FIG. 1a	Between about 6 mm and about 8 mm	6.5 mm
The sloping angle between the central section 1 and the "bottleneck" section 7a or 7b	"d" on FIG. 1a	Less than 45°	Less than 45°
Side of the core 5, in cross section**	"e" on FIG. 1b	Between about 0.5 mm and about 1.5 mm	1.0 mm
Diameter of the central section 1**	"f" on FIG. 1b	Between about 1.5 mm and about 2.5 mm	2.0 mm
Diameter of the end sections 2a and 2b**	"g" on FIG. 1b	Between about 6 mm and about 7 mm	6.0 mm

Comments. *The point 7e lies between about 0.05 mm and about 0.1 mm from the end face 4a or 4b, and the chamfer is formed between the point 7e and the end face. **This square and the two circles are substantially concentric, each having the center in the same point, with possible deviation for each center being within about 0.2 mm.

From the dimensions presented in Table 2 above, the dimension of the "bottleneck" section 7a or 7b can be easily calculated as being within a range of between about 3.8 mm and about 4.2 mm, preferably, about 4.0 mm.

Other limitations to the preferred structure of this invention comprise the following:

(a) the end faces 4a and 4b are tilted to between about 1° and about 3° , preferably, about 2° , from the perfectly vertical position, the end faces 4a and 4b being substantially parallel to each other, to within about 1° ;

(b) all diffusion bonded interfaces (e.g., between the inner surface of the cladding 6 and the outer surface of the core 5) are substantially free of gaps, voids, inclusions and like mechanical imperfections;

(c) the outer surface of the central section 1 is polished to reach a commercial grade degree of smoothness;

(d) all other outer surfaces, other than the end surfaces 4a and 4b, are polished to a rough grind of about 400 grit; and

(e) the end surfaces 4a and 4b are subjected to more thorough polishing and are then coated as described below.

The end surfaces 4a and 4b are polished to achieve a surface as provided by military specification MIL-O-1380. The surfaces 4a and 4b are then coated with the an anti-reflecting coating having properties in accordance with military specification MIL-C-48497, except for humidity requirements. Both military specifications are incorporated herein by reference.

The coating can be optionally, but not mandatorily, applied to the end chamfers. The coating must be able to withstand a flux density of at least 100 MW/cm².

The device creates a favorable thermal situation arising when the LuAG core absorbs the pump light and heats up internally. FIG. 2 is a graph of thermal analysis, showing the change in temperature of the active core (ΔT) as a function of the outer diameter of the cladding. The heat transfer coefficient of water cooling at the

surface of the cladding was also used as a variable parameter in order to illustrate advantages of the invention.

Because the LuAG core is hotter than the YAG cladding, and also because the LuAG linear thermal expansion coefficient is higher than that of the YAG cladding, the LuAG core expands relative to the YAG cladding. As a result, the LuAG core is under compressive stress during the pumping.

The above described situation, when the core is under compressive stress is preferred to the situation when the core is under the tensile stress (e.g., when the core shrinks relative to the cladding), when the core is subject to thermally induced failure, as the surrounding YAG cladding expands and destroys the LuAG core. The composition of the structure of this invention takes advantage of physical and thermomechanical properties of YAG and LuAG and avoids the extremely undesirable situation when the core is under tensile stress.

Having described the invention in connection with several embodiments thereof, modification will now suggest itself to those skilled in the art. As such, the invention is not to be limited to the described embodiments except as required by the appended claims.